ABSTRACT

Large variations exist between published mid-Cretaceous (late Barremian to early Turonian stages) seawater Sr-isotope stratigraphies; this has resulted in disparate interpretations of crustal production rates. We report on a detailed investigation of seawater Sr-isotope stratigraphy based on foraminifers and, where available, on inoceramid bivalves from 12 mid-Cretaceous Deep Sea Drilling Project and Ocean Drilling Program sections. The effects of diageneasis are assessed using scanning electron microscope observations and trace-elemental analyses, but are best distinguished by comparing the $^{87}\text{Sr}/^{86}\text{Sr}$ values of similar-age samples from different sites. Strontium-isotope analyses compiled from 9 of 12 sites that have detailed age control define one band of common values. This band is used as a composite curve, which presumably represents seawater $^{87}\text{Sr}/^{86}\text{Sr}$ values. The composite curve shows a “trough” of markedly lower $^{87}\text{Sr}/^{86}\text{Sr}$ values in the Aptian and early Albian stages, higher but constant values for the mid-Cretaceous (late Barremian to early Turonian stages) overlap with the pulse of mid-plate volcanic activity that produced the Ontong Java, Manihiki, and Kerguelen Plateaux. The exact age and the shape of the trough, however, are consistent with increased spreading rates at oceanic ridges, given the existing data on the timing of mid-plate volcanic activity.

INTRODUCTION

Major fluctuations in the strontium-isotope composition of seawater are recorded in Phanerozoic marine carbonate strata (e.g., Veizer and Compston, 1974; Brass, 1976; Burke et al., 1982; Hess et al., 1986; McArthur et al., 1993; Jones et al., 1994a). Because the ocean is well mixed with respect to Sr (the residence time of Sr in the oceans is on the order of $10^7$ yr and the mixing time of the oceans is on the order of $10^3$ yr), and because marine carbonates faithfully record oceanic Sr-isotope ratios at deposition (DePaolo and Ingram, 1985), stratigraphic fluctuations in marine carbonates are assumed to be globally synchronous. For the parts of the time scale where the seawater Sr-isotope curve is confidently established, it can be used to provide age control for other sedimentary rocks and to model geochemical cycles.

Significant advances have been made in the development of a Sr-isotope stratigraphy for the Neogene (e.g., DePaolo and Ingram, 1985; Hodell et al., 1989; Farrell et al., 1995). Because seawater $^{87}\text{Sr}/^{86}\text{Sr}$ values changed rapidly for much of this time interval, Sr isotopes can be used to date nonfossiliferous or poorly fossiliferous sediments with precision (e.g., Ludwig et al., 1988; McKenzie et al., 1988). Paleogene and Upper Cretaceous Sr-isotope stratigraphies are also well established (e.g., Denison et al., 1993; McArthur et al., 1993; Sugarman et al., 1995).

The Sr-isotope composition of seawater is controlled largely by variations in the input of Sr from continental weathering, hydrothermal circulation at spreading centers, and from carbonate dissolution. Therefore, changes in Sr-isotope ratios can elucidate the timing of orogenic events and the onset of glaciation (e.g., Raymo et al., 1988; Hodell et al., 1991), and has implications for the extent and timing of ridge-crest and mid-plate volcanism, and possibly even eustasy (e.g., Brass, 1976). Increasing Sr-isotope ratios generally indicate periods during which the supply of continental Sr ($^{87}\text{Sr}/^{86}\text{Sr} = 0.712$) increased with respect to the supply of Sr from mid-ocean ridges ($^{87}\text{Sr}/^{86}\text{Sr} = 0.703$), and conversely decreasing $^{87}\text{Sr}/^{86}\text{Sr}$ values suggest intervals when volcanic Sr increased with respect to Sr from the continents. Because the absence of magnetic reversals inhibits the dating of oceanic crust of mid-Cretaceous age, Sr-isotope curves have been used on both sides of the debate over the existence of rapid sea-floor spreading in this interval (e.g., Jones et al., 1994b; Ingram et al., 1994; Heller et al., 1996). High mid-Cretaceous crustal production rates at ridge crests and in mid-plate settings are inferred to have been associated with the ascent of a superplume from the mantle (e.g., Larson, 1991; Larson and Kincaid, 1996), a scenario with significant implications (e.g., Hays and Pitman, 1973). Despite the significance of these data, available Sr-isotope records for the mid-Cretaceous (late Barremian to early Turonian stages) possess great variability. The original studies of Burke et al. (1982) and Koepnick et al. (1985) show a significant amount of internal scatter, largely as a result of diagenetic alteration. Data from more recent studies show far less internal scatter. Pris-
tine remains of Cenomanian-Turonian inoceramids and ammonites from the Western Interior basin of North America were used in the Sr-isotope study of McArthur et al. (1994). Jones et al. (1994b) measured $^{87}$Sr/$^{86}$Sr values of original belemnites and oysters obtained from sequences in England, and Ingram et al. (1994) measured Sr-isotope ratios on apatitic remains of fish teeth concentrated from mid-Cretaceous limestone sequences in Italy. Strontium-isotope values of inoceramids from the Albian section of Deep Sea Drilling Project (DSDP) Site 511 on the Falkland Plateau in the South Atlantic were reported by Huber et al. (1995). Jenkyns et al. (1995) measured $^{87}$Sr/$^{86}$Sr values on Hauterivian-Albian shallow-water carbonates from Ocean Drilling Program (ODP) Site 866 atop Resolution Guyot in the central Pacific. Large disparities continue to exist between these recent papers. In particular, the Sr-isotope ratios of Ingram et al. (1994) are noticeably higher than those reported by other authors.

The disparities among current mid-Cretaceous Sr-isotope stratigraphies may result from modification of original isotope ratios by diagenetic alteration, or analytical procedures that have sampled Sr from nonbiogenic sources such as clays. Errors also exist in establishing the ages of the samples that come from a geographically diverse set of locations. For example, it is often difficult to correlate precisely between Tethyan and higher latitude sequences, or between sediments deposited in pelagic and shallower water environments.

We present an independent mid-Cretaceous seawater Sr-isotope stratigraphy based on measurements of foraminifers and inoceramids from 12 DSDP and ODP sections. The deep-sea record offers advantages for establishing Sr-isotope stratigraphies: (1) sections are usually more complete than their shallower water counterparts; and (2) cosmopolitan faunal and floral assemblages provide more direct correlation with major tectonic and paleoceanographic events, enabling a thorough analysis of the causes of changes in seawater $^{87}$Sr/$^{86}$Sr values.

**SITES, SECTIONS, AND METHODS**

**Sites and Sections**

The 12 sites investigated are distributed throughout the major ocean basins (Fig. 1) and include pelagic and hemipelagic depositional settings characterized by nannofossil-rich claystone and chalk (Table 1). The sampled sites compose an expanded, largely continuous, and well-recovered upper Barremian to lower Turonian sequence. Sample spacing averages 1.5 m, equivalent to a temporal duration of about 150000–250000 yr at standard sedimentation rates for pelagic carbonates.

**Time Control and Correlation**

Establishment of Ages and Biostratigraphic Error. Calcareous nannofossil biostratigraphic control exists in all of the sections analyzed and many are also constrained by planktic foraminiferal biostratigraphy (Table 1). Only one site has magnetostratigraphy: basal Aptian magnetic polarity zone M0 was identified at Site 641 (Ogg, 1988).

Chronological ages of all samples are estimated using the integrated calcareous nannofossil-planktic foraminiferal biostratigraphic scheme of Bralower et al. (1993) calibrated to the time scale of Gradstein et al. (1994). The calibration generally follows Premoli Silva and Sliter (1994) and Erba et al. (1996), who calculated planktic foraminiferal and calcareous nannofossil zonal ages from their relative positions in sections from the Umbrian Apennines of Italy assuming constant sedimentation rates. The ages of all datum levels are shown in Figure 2.

Ages of samples in Sites 258, 511, 545, 547, 551, 641 and 763 are estimated by assuming constant sedimentation rates between a series of planktic foraminiferal and nannofossil datum levels (Table 2). We use the most reliable datum levels in each section. Chronological ages of samples from sections that are incomplete or poorly recovered are more difficult to determine. At Sites 390, 392, 417, and 418, only one datum...
Sample 5-2, 101 cm, lies in the uppermost part of the Globigerinelloides algerianus zone. The FO of Ticinella primula lies between samples 4-CC and 5R-1, 88 cm. Between samples 3-1, 94 cm, and 3-2, 102 cm, the nannofossil E. floralis, but absence of the foraminifers G. algerianus and G. ferreolensis. Core 17 lies in the upper Albian Rotalipora ticinensis zone based on the presence of R. ticinensis, and absence of R. cushmani, "Globigerinelloides" gyroidinaeformis, specimens belonging to the G. blowi group, the nannofossils Eprolithus floralis and Parahabdolithus achlyostaurion, and the absence of the foraminifers Globigerinelloides algerianus, G. ferreolensis, and G. aptiense, and the nannofossil Prediscosphaera columnata.

<table>
<thead>
<tr>
<th>Site</th>
<th>Location, Interval</th>
<th>Lithology</th>
<th>Biostratigraphy</th>
</tr>
</thead>
<tbody>
<tr>
<td>DSDP 258</td>
<td>Naturaliste Plateau, Indian Ocean</td>
<td>Light olive-gray nannofossil chalk</td>
<td>Thierstein (1974)(n)</td>
</tr>
<tr>
<td>DSDP 327</td>
<td>Falkland Plateau, South Atlantic</td>
<td>Pink nannofossil chalk</td>
<td>Wise and Wind (1977)(n)</td>
</tr>
<tr>
<td>DSDP 390</td>
<td>Blake Nose, North Atlantic</td>
<td>Gray-yellow marly nannofossil ooze</td>
<td>Gradstein (1979)(f); Schmidt (1979)(n); this study (n,f)</td>
</tr>
<tr>
<td>DSDP 392</td>
<td>Blake Nose, North Atlantic</td>
<td>Variable: Light gray-yellow-pink marly nannofossil ooze</td>
<td>Gradstein (1979)(f); Schmidt (1979)(n); this study (n,f)</td>
</tr>
<tr>
<td>DSDP 417</td>
<td>Bermuda Rise, North Atlantic</td>
<td>Variable: Light gray-brown-pink marly nannofossil ooze</td>
<td>Miles and Orr (1980)(f); Gartner (1980)(n); Siesser (1980)(n); this study (n,f)</td>
</tr>
<tr>
<td>DSDP 418</td>
<td>Bermuda Rise, North Atlantic</td>
<td>Variable: Pink marly chalk and light gray claystone</td>
<td>Miles and Orr (1980)(f); Gartner (1980)(n); Siesser (1980)(n); this study (n,f)</td>
</tr>
<tr>
<td>DSDP 511</td>
<td>Falkland Plateau, South Atlantic</td>
<td>Pink nannofossil chalk</td>
<td>Krashenninikov and Basov (1983)(f); Bralower et al. (1993)(f); Wise (1983)(n); Bralower (1992)(n); this study (f)</td>
</tr>
<tr>
<td>DSDP 545</td>
<td>Morocco Basin, North Atlantic</td>
<td>Green nannofossil claystone</td>
<td>Leckie (1984)(f); Wiegand (1984)(n); Bralower (1992)(n); this study (f)</td>
</tr>
<tr>
<td>DSDP 547</td>
<td>Morocco Basin, North Atlantic</td>
<td>Green nannofossil claystone</td>
<td>Leckie (1984)(f); Wiegand (1984)(n); Bralower (1992)(n); this study (f)</td>
</tr>
<tr>
<td>DSDP 551</td>
<td>Goban Spur, North Atlantic</td>
<td>White nannofossil chalk</td>
<td>Graciansky et al. (1985)(f); Bralower (1988)(n)</td>
</tr>
<tr>
<td>ODP 641</td>
<td>Galicia Bank, North Atlantic</td>
<td>Gray nannofossil claystone</td>
<td>Applegate and Bergen (1986)(n); Bralower et al. (1994)(n)</td>
</tr>
<tr>
<td>ODP 763</td>
<td>Exmouth Plateau, Indian Ocean</td>
<td>Light green zeolitic nannofossil claystone</td>
<td>Bralower and Siesser (1992)(n); this study (f)</td>
</tr>
</tbody>
</table>

Note: DSDP—Deep Sea Drilling Project, ODP—Ocean Drilling Program, (n)—nannofossils, (f)—foraminifera.
Figure 2. The chronostratigraphic scheme utilized in this study. Integrated nannofossil and planktic foraminiferal zonation is after Bralower et al. (1993), calibrated to the time scale of Gradstein et al. (1994), following Premoli Silva and Sliter (1994) and Erba et al. (1996). Ages of the Aptian nannofossil subzonal datum levels are estimated directly from their correlation with foraminiferal datum levels or stage boundaries where they correlate closely. In the Albian stage, where the sections studied by Premoli Silva and Sliter (1994) appear to be condensed, the ages of nannofossil subzonal datum levels are calculated from their relative positions between the first occurrences of *Prediscosphaera columnata* and *Eiffellithus turriseiffelii* in the expanded and apparently complete Albian section at Ocean Drilling Program Site 763 (from Bralower and Siesser, 1992), assuming constant sedimentation rates.
can be accurately determined in any continuous stratigraphic interval. Here a constant sedimentation rate of 5 m/m.y. above and below this datum is assumed, a rate typical for Holocene calcareous oozes from the North Atlantic (Berger, 1974). The major source of error in determining the ages of samples is associated with the assumption of constant sedimentation rates. This error is significantly larger in sections that are incompletely recovered, because datum levels cannot be placed precisely. In these cases the magnitude of the potential error approaches the duration of the zonal unit of interest, although we believe that the errors are much less than this.

**Correlation with Previous Sr-Isotope Stratigraphies.** Our calculated sample ages enable correlation with previously published Sr-isotope stratigraphies based on macrofossil biostratigraphy (McArthur et al., 1994; Jones et al., 1994b) after these data sets were recalibrated to the Gradstein et al. (1994) time scale. The sections investigated by Ingram et al. (1994) have planktic foraminiferal biostratigraphic control allowing direct comparison with the chronostratigraphic scheme derived here (Fig. 2). Samples analyzed by Ingram et al. (1994) were calibrated to the Harland et al. (1989) time scale, but the data of Koepnick et al. (1985), which they incorporated, appear to have remained correlated to the Harland et al. (1982) time scale. In addition, the biostratigraphy of samples from Site 167 in Koepnick et al. (1985) was taken from Winterer et al. (1973) and modifications to the biostratigraphy of this site by Tarduno et al. (1989) resulted in changes in chronological ages of as much as 10 m.y. To compare the Ingram et al. (1994) stratigraphy with other data, we have recalibrated the Koepnick et al. (1985) ages to the Gradstein et al. (1994) time scale by (1) using the revised planktic foraminifer biochronology of Site 167 from Tarduno et al. (1989), and (2) assuming similar relative stage positions for the ages given by Koepnick et al. (1985) for the Iranian samples.

**Sample Preparation**

Although different processing techniques are used to extract foraminifers for Sr-isotope analyses (e.g., Hodell et al., 1989; DePaolo and Finger, 1991; Sugarman et al., 1995), we avoided use of chemicals such as Calgon and kerosene to break up material because of uncertain effects on the Sr-isotope ratios. About 5 cm³ of material was broken into pea-sized pieces in a pestle and mortar. These pieces were soaked in kerosene to break up material because of unavoidable use of chemicals such as Calgon and kerosene to break up material. The samples were washed onto filter papers and dried in a 60°C oven.

Foraminifers were picked for isotope analysis using a binocular microscope. No distinct was made between planktic and benthic foraminifers. Where available, approximately 50 of the best preserved specimens regardless of taxonomic category were picked from the >250 µm fraction; if this size fraction contained an inadequate number of foraminifers, as many as 150 specimens were separated from the 125–250 µm fraction. Only in the upper Barremian–lower Aptian interval of Site 641 was it necessary to pick the foraminifers from the 63–125 µm fraction. Where other fossil material was present, e.g., ostracodes or inoceramids, several fragments were isolated for separate Sr-isotope measurements. A few bulk carbonate Sr-isotope analyses were carried out at Site 763.

The degrees of overgrowth and infilling of foraminifers were noted in each sample. To document preservation further, two typical foraminiferal specimens from each sample were observed in a Leica Stereoscan 440 scanning electron microscope (SEM) at the University of North Carolina–Chapel Hill. Several samples of variable external preservation were broken open to observe the interior of the chambers. Energy dispersive X-ray spectrometry was used to identify nongenetic mineral phases.

**Strontium-Isotope Analysis**

Samples were dissolved in 500 µL of 1 M acetic acid, centrifuged, and the decantate was dried. The sample was then dissolved in 250 ml of 5 N HNO₃, loaded on a column containing 50 µl of Eichrom SrSpec resin, washed with 750 µl of 5 N HNO₃, and the Sr was collected in 1 ml of H₂O. Total procedural blanks were 100 to 200 pg. Small samples (0.2–1 mg) were loaded in TaCl₃ on single Re filaments; larger samples (1–5 mg) were loaded on single Ta filaments. Samples were analyzed at the University of North Carolina–Chapel Hill on a VG Sector 54 thermal ionization mass spectrometer in dynamic mode. Fractionation is corrected using ⁸⁶Sr/⁸⁸Sr = 0.1194. Strontium-isotope values are archived in Table DR1.

In this study, ⁸⁷Sr/⁸⁶Sr analyses of 330 aliquots of Sr carbonate standard SRM987 yielded an average value of 0.710246 ± 3 (2σ). Part of this uncertainty is the result of degradation of the collectors used to measure the relative intensity of the Sr ion beams. In fact, some collectors were replaced twice during the course of this study. To minimize instrumental variation, we analyzed three to six, usually five, SRM987 aliquots in each sample carousel, and used the values obtained to adjust the ⁸⁷Sr/⁸⁶Sr ratios of the samples. We accepted 0.710250 as the correct ⁸⁷Sr/⁸⁶Sr value of SRM987. If, for example, the values for the standards in a particular turret averaged 0.710246, we added 0.000004 to the ⁸⁷Sr/⁸⁶Sr ratio for each sample. This procedure reduced the uncertainty for all 330 analyses of SRM987 to 625 (2σ). We analyzed 28 samples in duplicate (see footnote 1); for 90% of these samples, the individual ⁸⁷Sr/⁸⁶Sr ratios were within ±15 of the mean of the duplicate analyses. Internal precision for Sr-isotope analysis was typically 0.0006% to 0.0009% standard error, based on 100 dynamic cycles of data collection.

Tests were conducted to estimate variations in ⁸⁷Sr/⁸⁶Sr values of the utilized fossil materials. Different foraminiferal fractions from the same sample (combined planktics and benthics of all size fractions, combined planktics and benthics >250 µm, separate planktics, separate benthics) show negligible differences in ⁸⁷Sr/⁸⁶Sr values (Fig. 3), justifying our measurement of mixed foraminifers.
Trace Elemental Analysis

To evaluate further the extent of postdepositional alteration of the samples, we determined trace- and minor-element:calcium ratios on representative splits of foraminifers and inoceramids prisms (where present) from carefully selected samples from all sites except Site 258. Samples were dissolved in 0.5 N nitric acid and analyzed for their Ca, Mg, Sr, Na, Fe, Mn, and Ba content on a Fisons Spectraspan 7 direct current plasma atomic emission spectrophotometer at the Department of Geology, Duke University. Matrix-matched instrument-calibration standards were prepared from Spex brand pure, single-element plasma-grade solutions. An internal consistency standard prepared from an in-house limestone standard (PE3) and plasma-grade solutions was analyzed repeatedly to monitor run-to-run precision. On the basis of these repeated analyses of standard PE3, precision for Mg/Ca, Mn/Ca, and Ba/Ca ratios is better than ±4% (2σ), whereas precision for Na/Ca and Fe/Ca ratios is better than ±8%. Trace-elemental data are compiled in Table DR1 (see footnote 1).

RESULTS

Foraminiferal Preservation

Light microscope observations indicate that foraminifer preservation is variable between sites and within individual sections. Scanning electronic microscope (SEM) inspection of the wall texture and chamber filling suggests that, in many cases, light microscopic observations are unsuitable for evaluating the diagenetic alteration of the foraminifers analyzed. For example, specimens from Site 547 which possess a nearly pristine wall texture and appear translucent in the light microscope are filled with sparry calcite or other authigenic mineral phases. Conversely, foraminifers from Site 641 that appear to be poorly preserved in the light microscope are nearly hollow and have only slightly overgrown wall textures. Foraminiferal preservation in the sites investigated is summarized in Table 3. Figure 4 illustrates SEM micrographs of specimens showing the complete range of preservation observed. More detailed illustrations of foraminiferal preservation are given in Fassell and Bralower (in press).

Strontium-Isotope Data

Site 511. Strontium-isotope ratios of foraminifers and inoceramids show similar stratigraphic trends for most of the section studied at Site 511 (Fig. 5). Foraminiferal 87Sr/86Sr values increase from close to 0.707250 near the base of nannofossil zone NC8 (480 m below sea floor [mbsf]) to close to 0.707450 at the base of zone NC9 (445 mbsf), and decrease slightly in zones NC9 and NC10 (435 mbsf) (Fig. 5). Strontium-isotope ratios measured on inoceramids have slightly less inter-sample variability, and have average values that are 0.000016 lower than the foraminifers. Between 493 and 481 mbsf, trends are different, and inoceramid record values of as much as 0.0001 lower (Fig. 5).

Site 545. Considerable intersample variability exists in 87Sr/86Sr values in zone NC7 and the lower part of zone NC8 (420–530 mbsf) (Fig. 6). Values appear to decrease upward from 0.70745 at 530 mbsf to 0.7073 at 515 mbsf. Strontium-isotope values fluctuate around 0.7073 from 515 to 420 mbsf. Values increase markedly between 390 and 375 mbsf across the unconformity between zones NC8 and NC9; the abrupt increase apparently results from mixing of foraminifers of different ages in a slump in core 40. Thus Sr-isotope measurements from this core were excluded from subsequent analysis. Strontium-isotope values remain close to 0.70745 and show less variability in zone NC10 and the lower part of zone NC11 (360 to 260 mbsf).

Site 547. The 87Sr/86Sr values of foraminifers at Site 547 show few consistent long-term trends (Fig. 7). Samples between 780 and 740 mbsf tend to have higher values (approximately 0.70755) than those above 700 mbsf, where values fluctuate between 0.70730 and 0.70755.

Hole 763B. The biostratigraphically complete upper Aptian to lower Turonian section at Site 763 records no long-term trends in 87Sr/86Sr values measured on foraminifers (Fig. 8). Instead, this section shows 2–20 m 0.0001–0.0003 amplitude, multisample fluctuations. Bulk and inoceramid 87Sr/86Sr values are predominantly higher than those measured on foraminifers in the same samples.

Other Sites. Strontium-isotope measurements of samples from Sites 258, 327, 390, 392, 417, 418, 551, and 641 are illustrated in Figures 9–16. In general, considerably less intersample variability is observed within individual nannofossil zones than between zones. Inoceramid 87Sr/86Sr values are slightly lower than those measured on foraminifers (Figs. 10 and 12).

DISCUSSION

Monitoring Diagenesis of Samples

The Sr-isotope composition of pore waters of marine sediments may be different from the carbonates with which they are associated (Elderfield and Gieskes, 1982; Richter and DePaolo, 1988; Paull et al., 1995), and recrystallization may lead to altered Sr-isotope ratios. However, the presence of secondary calcite does not necessarily signify diagenetically altered Sr-isotope ratios, because the Sr in the diagenetic calcite may be obtained from local dissolution (e.g., Jenkyns et al., 1995). Even small amounts
Figure 4. (a) Sample 511-51-6, 12–14 cm, wall structure shows unfilled pores. (b) Sample 551-5-1, 91–94 cm, wall structure shows minor overgrowth. (c) Sample 392A-3-1, 86–88 cm, wall shows minor overgrowth on inside, but little pore infilling. (d) Sample 392A-2-2, 109–112 cm, wall shows moderate overgrowth on inside. (e) Sample 641C-10R-1, 87–89 cm, broken specimen shows rhombs of calcite growing inside chamber. (f) Sample 511-55-2, 15–17 cm, moderately overgrown wall texture. (g) Sample 327A-21-3, 133–136 cm, inside of a chamber shows calcite infilling. (h) Sample 641C-11R-2, 3–6 cm, broken planktic specimen shows overgrown outer wall and rhomb of calcite lodged inside. (i) Sample 763B-35X-5, 33–40 cm, broken specimen shows slightly overgrown wall and chamber infilling of zeolite (large blady crystals), coccoliths, and minute cristobalite lepispheres. (j, k) Sample 547A-54-2, 26–29 cm, broken specimen shows entirely infilled chamber and overgrown spines in photo k. (l) Sample 547A-45-2, 34–37 cm, broken specimen of Hedbergella sp. shows entirely infilled chamber, pristine wall and incipient pore infilling. Scale bars represent 5 µm. See Table 3 for summary of preservation of sites.
**Figure 5.** Strontium-isotope data from Site 511. Nannofossil biostratigraphy is after Wise (1983), modified by Bralower (1992). Planktic foraminiferal biostratigraphy is modified after Bralower et al. (1993). An age model is shown on the left; arrows signify tie points (see Table 2). The base of core 49 is placed in the upper Cenomanian stage, following Krasheninnikov and Basov (1983). Cen—Cenomanian, mbsf—meters below sea floor.

**Figure 6.** Strontium-isotope data from Site 545. The nannofossil biostratigraphy is after Wiegand (1984), modified by Bralower (1992). The planktic foraminiferal biostratigraphy is modified after Leckie (1984). The position of the slump is indicated. An age model is shown on the left; arrows signify tie points (see Table 2). The key to the lithologic column is given in Figure 5. mbsf—meters below sea floor.
Figure 7. Strontium-isotope analyses of foraminifers from Site 547. Nannofossil biostratigraphy is after Wiegand (1984), modified by Bralower (1992). The planktic foraminiferal biostratigraphy is after Leckie (1984). An age model is shown on the left; arrows signify tie points (see Table 2). The key to the lithologic column is given in Figure 5. mbsf—meters below sea floor.

Figure 8. Strontium-isotope data from Hole 763B. The nannofossil biostratigraphy is after Bralower and Siesser (1992). Planktic foraminiferal biostratigraphy is from this study. An age model is shown on the left; arrows signify tie points (see Table 2). The key to the lithologic column is given in Figure 5. Tur.—Turonian, mbsf—meters below sea floor.
Figure 9. Strontium-isotope analyses of foraminifers from Site 258. The nannofossil biostratigraphy is after Thierstein (1974). An age model is shown on the left; arrows signify tie points (see Table 2). The key to the lithologic column is given in Figure 5. Turon.—Turonian, mbsf—meters below sea floor.

Figure 10. Strontium-isotope data from Hole 327A. The nannofossil biostratigraphy is after Wise and Wind (1977). An age model is shown on the left; the arrow signifies tie point (see Table 2). The key to the lithologic column is given in Figure 5. mbsf—meters below sea floor.
Figure 11. Strontium-isotope analyses of foraminifers from Site 390. The nannofossil biostratigraphy is after Schmidt (1979), modified as part of this study. Planktic foraminiferal biostratigraphy is modified after Gradstein (1978). An age model is shown on the left; arrows signify tie points (see Table 2). The key to the lithologic column is given in Figure 5. Camp.—Campanian, mbsf—meters below sea floor.

Figure 12. Strontium-isotope data from Hole 392A. The nannofossil biostratigraphy is after Schmidt (1979), modified as part of this study. Planktic foraminiferal biostratigraphy is modified after Gradstein (1978). An age model is shown on the left; arrows signify tie points (see Table 2). The key to the lithologic column is given in Figure 5; mbsf—meters below sea floor.

Figure 13. Strontium-isotope analyses of foraminifers from Hole 417D. The nannofossil biostratigraphy is after Gartner (1980) and Siesser (1980). Foraminiferal biostratigraphy is modified after Miles and Orr (1980). An age model is shown on the left; arrows signify tie points (see Table 2). The key to the lithologic column is given in Figure 5; mbsf—meters below sea floor.

Figure 14. Strontium-isotope analyses of foraminifers from Hole 418B. The nannofossil biostratigraphy is after Gartner (1980) and Siesser (1980). Foraminiferal biostratigraphy is modified after Miles and Orr (1980). An age model is shown on the left; the arrow signifies tie point and crosses indicating interpreted ages of samples (see Table 2 and section 2.2). The key to the lithologic column is given in Figure 5; mbsf—meters below sea floor.
of clay in a sample can increase the apparent \(^{87}\text{Sr}/^{86}\text{Sr}\) values of carbonates, because some of this Sr could be extracted during the preparation of the sample (e.g., Dasch and Biscaye, 1971; Burke et al., 1982). We have attempted to remove adhering clay from particles prior to analysis, and the dilute (1 M) acetic acid used to dissolve samples minimizes the effect from any remaining clay. In all of the chalk samples analyzed, very little or no clay residue was observed, and such contamination seems unlikely. However, some Sr extraction could have occurred in claystone samples from Sites 545, 547, and 763 where light green clay residues reappeared in claystone samples from Sites 545, 547, and 548. However, some Sr extraction could have occurred in claystone samples from Sites 545, 547, and 763 where light green clay residues remained in many samples after dissolution.

**Trace-Element and Preservational Evidence.** SEM observations show some overgrowths of minute rhombs on the inside of chamber walls in almost all foraminiferal samples analyzed, regardless of their pristine appearance (Fig. 4). Volumes of these minute rhombs are negligible compared to the volume of primary calcite in samples from the upper part of Site 511, and Sites 258, 390, 392, 417, and 418. However, in other samples, such as those from the lowermost part of Site 511, and most of those analyzed at Sites 545, 547, and 641, pores and whole chambers are infilled with secondary calcite (Table 3; Fig. 4).

Trace-elemental analyses help identify samples with potentially altered Sr-isotope ratios. Diagenetic calcite tends to have lower Sr/Ca ratios than biogenic calcite (e.g., Baker et al., 1989). Ratios of Na/Ca show the same pattern, whereas Fe/Ca and Mn/Ca ratios in diagenetic calcite may be elevated compared to marine biogenic calcite (e.g., Veizer, 1983). Interpretation of trace-elemental analyses of foraminifers is difficult, however, because a contribution of these elements may be derived from noncalcitic phases coating the outside of tests and filling chambers. Certain phases, such as pyrite and Fe-Mn oxides, may contribute no Sr; others, such as clays and zeolites, may have high Sr contents, at least part of which may be leached by dilute acids during sample preparation. Because of these problems, we use trace-elemental analysis to identify diagenetically altered \(^{87}\text{Sr}/^{86}\text{Sr}\) values in a site-by-site basis.

Ratios of Sr/Ca, Na/Ca, Fe/Ca, Mn/Ca, and Ba/Ca of all foraminiferal samples analyzed are plotted versus age in Figure 17. These plots show significant differences between sites. We believe that these differences result from (1) contamination with Sr-bearing phases such as clays and zeolites in some sites, and non-Sr bearing phases such as Fe-Mn oxides in others, and (2) differences in the amount of calcite recrystallization.

High Sr/Ca, Na/Ca, and Ba/Ca ratios in most samples analyzed at Site 763 (Fig. 17) are thought to result from the presence of zeolites observed growing inside foraminifer tests (Fig. 4i). Strontium-isotope values substantially higher than those in contemporaneous sections (Fig. 18) probably result from the growth of these zeolites from fluids with higher \(^{87}\text{Sr}/^{86}\text{Sr}\) ratios. We note that some of the Sr-isotope fluctuations at Site 763 appear to parallel those at other sites. For example, increases in \(^{87}\text{Sr}/^{86}\text{Sr}\) values in the lowermost Albian stage (between 112 and 110 Ma) correspond to an increase at Site 511 in this same time interval (Fig. 18). This suggests a systematic addition of high-ratio Sr to each sample. Most foraminifers from Site 547 are nearly filled with calcite (Fig. 4; j–l), and relatively low Sr/Ca and Na/Ca ratios (Fig. 17) are additional indicators of diagenetic alteration. Trace-elemental analyses can also be used to interpret Sr-isotope records with large amounts of variability such as the Aptian part of the record at Site 545 (Fig. 6). Here there is a significant correlation between fluctuations in \(^{87}\text{Sr}/^{86}\text{Sr}\) values, and Sr/Ca and Na/Ca ratios (Fig. 19). Samples with higher \(^{87}\text{Sr}/^{86}\text{Sr}\) values generally possess lower Sr/Ca and Na/Ca ratios, indicating a larger proportion of secondary calcite. In the interval between 390 and 510 mbsf, therefore,
only the samples with the highest Sr/Ca (>0.9) and Na/Ca ratios (>~8) are interpreted as reliable carriers of original Sr-isotope ratios. Strontium-isotope values from the Albian and Cenomanian of Site 545 (390 to 255 mbsf) show much less scatter (Fig. 18) and samples in this interval have generally higher Sr/Ca ratios (Fig. 17). The trace-elemental data suggest a lower degree of recrystallization or a lower clay content, and the Albian and Cenomanian $^{87}\text{Sr}/^{86}\text{Sr}$ values from Site 545 are thought to be reliable monitors of past seawater.

Comparison of trace-elemental analyses of foraminifers and inoceramids in samples from Site 511 provides an opportunity to evaluate the effect of diagenesis on $^{87}\text{Sr}/^{86}\text{Sr}$ values. Above 481 mbsf, where preservation of both groups is excellent, $^{87}\text{Sr}/^{86}\text{Sr}$ values lie largely within error limits (Fig. 5), suggesting a primary seawater record. Below this depth, infilling of foraminiferal chambers increases and $^{87}\text{Sr}/^{86}\text{Sr}$ values progressively diverge from those of inoceramids. Alteration of foraminiferal $^{87}\text{Sr}/^{86}\text{Sr}$ values is indicated by increasing Mn/Ca ratios, generally lower Sr/Ca values (Figs. 17 and 20) and anomalously low $\delta^{18}O$ values (Fassell and Bralower, in press). Because the Site 511 inoceramid $^{87}\text{Sr}/^{86}\text{Sr}$ values show less variability, and are generally slightly lower than those of foraminifers (Figs. 3 and 5), and because inoceramids possess more uniform trace-element contents (Fig. 20), we utilize them in developing the seawater Sr-isotope curve.

Stacking of $^{87}\text{Sr}/^{86}\text{Sr}$ Values from Different Sites and Construction of the Composite Record. When all of the $^{87}\text{Sr}/^{86}\text{Sr}$ values of contemporaneous samples from different locations are stacked (Fig. 18), a significant amount of scatter exists. However, when all of the samples with indications of diagenetic alteration or suspect age are removed, a much narrower band at the lower limits of the total range is defined (Fig. 21). We maintain that this band represents the best record of seawater Sr-isotope variation and use the $^{87}\text{Sr}/^{86}\text{Sr}$ values in this common band.
to define our composite record. The 150 data points used in the composite record include: (1) all measurements of foraminifers from Sites 258, 390, 417, 418, 551, and 641, (2) all inoceramid analyses from Sites 327, 392, and 511 (and foraminiferal measurements in the former two sites in samples in which no inoceramid analyses were performed), and (3) Cenomanian, Albian, and certain Aptian data points from Site 545 (see above discussion). Strontium-isotope values of some 230 samples, including many from Site 545, and most from Sites 547 and 763 lie significantly above this band (Fig. 18). We maintain that the $^{87}\text{Sr}/^{86}\text{Sr}$ values of these samples have increased during diagenesis.

**Deep-Sea Sr-Isotope Stratigraphy**

The composite record shows a decrease in $^{87}\text{Sr}/^{86}\text{Sr}$ values from just over 0.70745 in the late Barremian stage to just over 0.70720 in the late Aptian stage (Fig. 21). Strontium-isotope values begin to increase close to the Aptian-Albian boundary, rise rapidly in the early Albian and remain close to 0.70745 from the middle Albian to the late Cenomanian stages. A sharp decrease in $^{87}\text{Sr}/^{86}\text{Sr}$ values occurs just above the Cenomanian-Turonian boundary; earliest Turonian values average 0.70735. The curve produced by local linear smoothing (hereafter referred to as the smoothed composite curve) accounts for most of the variation shown by the composite data set (Fig. 22A). One exception is in the latest
Aptian (Ticinella bejaouaensis foraminiferal zone), where a short-term increase in $^{87}\text{Sr}/^{86}\text{Sr}$ values is seen in Sites 392, 418, and 545, but not in Sites 417 and 511. Although we believe that the lower $^{87}\text{Sr}/^{86}\text{Sr}$ values are more reliable, and thus the trough in the late Aptian stage may actually be slightly larger than the smoothed composite curve indicates, the possibility of two smaller troughs with a minor saddle in the $T$. bejaouaensis zone cannot be dismissed.

Comparison of our composite deep-sea Sr-isotope record with published data sets shows both similarities and differences. The smoothed composite curve shows a close resemblance to the values of McArthur et al. (1994) for the Cenomanian–early Turonian (Fig. 22E). The composite data show significantly less scatter than those of Ingram et al. (1994) (Fig. 22, A and B). We have smoothed the Sr-isotope ratios of Ingram et al. (1994), including both original and revised (Tarduno et al., 1989) biostratigraphic ages of the samples of Koepnick et al. (1985) from Site 167, and make the following observations: (1) smooths from the corrected and uncorrected data are approximately parallel to the smoothed composite curve, but $^{87}\text{Sr}/^{86}\text{Sr}$ values of Ingram et al. (1994) are higher by an average of 0.0001–0.00015 (Fig. 22B); (2) smoothed curves drawn through the Ingram et al. (1994) data set show a considerably smaller Aptian-Albian trough than the curve drawn by Ingram et al. (1994); and (3) if two age points at 111.4 and 112.6 Ma are excluded, the trough in the smoothed curve of the Ingram et al. (1994) data set virtually disappears. Thus we conclude that the curve of Ingram et al. (1994) is arbitrarily drawn. We postulate that the differences between the Ingram et al. (1994) data set and the composite result from addition of Sr with relatively high $^{87}\text{Sr}/^{86}\text{Sr}$ values during diagenesis, and/or from the difficulty of removing particles such as clay with high $^{87}\text{Sr}/^{86}\text{Sr}$ values from the fish teeth.

The shape of our composite curve is comparable to that defined by the Barremian-Albian data of Jones et al. (1994b); however, it is clear that there are consistent age offsets of as much as 4 m.y. (Fig. 22C). The analyses of Jones et al. (1994b) derive from well-preserved clams and oysters in shallow-water sections from England. Ages in these sections were determined using Boreal ammonite-subzonal and stage-boundary definitions assuming equal subzone durations. We have used existing nannofossil zonal-ammonite subzonal correlations (Table 4) to re-calibrate the Jones et al. (1994b) data set and produce a modified curve by smoothing. The resulting curve lies remarkably close to the smoothed composite curve (Fig. 22D). The apparently larger scatter of the composite compared to the Jones et al. (1994b) and McArthur et al. (1994) data sets is likely a result of (1) the significantly larger size of our data set, and (2) the longer durations of the microfossil zonal units than ammonite subzones.

The composite Sr-isotope record shows remarkable similarity to the curve of Jenkyns et al. (1995) obtained from the analysis of shallow-water limestones recovered in ODP Site 866 from Resolution Guyot in the central Pacific (Fig. 22F). This independent data set provides additional support for the composite Sr-isotope record.

The steeper portions of the smoothed composite curve offer significant chronologic potential, especially for unfossiliferous sedimentary rocks. We offer polynomial and linear equations fitting three portions of the smoothed composite curve (Table 5).

Our data allow correlation of Sr-isotope values with microfossil zonal boundaries, and thus modification of previous correlations of the Sr-isotope stratigraphy and stage boundaries. Two boundaries lie at diagnostic points in the composite curve: the Aptian-Albian boundary is close to the minimum $^{87}\text{Sr}/^{86}\text{Sr}$ value in the broad Aptian–early Albian $^{87}\text{Sr}/^{86}\text{Sr}$ trough, and $^{87}\text{Sr}/^{86}\text{Sr}$ values decrease rapidly just above the Cenomanian-Turonian boundary (Figs. 21 and 22A). Therefore the composite Sr-isotope stratigraphy has potential for helping to identify these stage boundaries.

Geologic Significance of Mid-Cretaceous Sr-Isotope Curves

Case for High Crustal Production Rates. Larson (1991) and Larson and Olson (1991) proposed that the volume of mid-plate volcanic plateaus formed in the mid-Cretaceous was almost three times greater and that production of ridge-crest crust was 28% higher than in prior and subsequent time intervals. Coincidence of high mid-plate and ridge-crest crustal production rates is thought to have resulted from a mantle plume that fueled volcanism in both settings (e.g., Larson and Kincaid, 1996).

Isotopic, biostratigraphic, and magnetostratigraphic age constraints of mid-Cretaceous mid-plate and ridge-crest crust are few, however. For example, a major episode of flood basalt volcanism occurred in the late Barremian to early Aptian stages when the Hess Rise and the Ontong Java and Manihiki Plateaus in the central and western Pacific were constructed (Schlanger et al., 1981; Duncan and Richards, 1990; Tarduno et al., 1991). The age of the largest and best dated of these features, the Ontong Java Plateau, is constrained by isotopic dates on basalt from only four locations. These dates range between 120 and 124 Ma, but cluster between 121.5 and

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Figure 19. Sr/Ca and Na/Ca ratios (µmol/mol) plotted vs. $^{87}\text{Sr}/^{86}\text{Sr}$ values of samples in the Aptian section of Site 545. Pearson correlation coefficients for $^{87}\text{Sr}/^{86}\text{Sr}$ values vs. Sr/Ca and Na/Ca are –0.53 and –0.43, respectively which are both significant at the 99% interval of confidence. See text for discussion.
Figure 20. Elemental ratios (µmol/mol) plotted vs. $^{87}\text{Sr}/^{86}\text{Sr}$ values of samples from Site 511. Open squares refer to foraminiferal analyses from above 481 m below the sea floor (mbsf) (section 511-55-1 and up); closed squares refer to foraminiferal analyses from below 481 mbsf (section 511-55-2 and down); triangles refer to inoceramid analyses.
Figure 21. Composite Sr-isotope data set. All samples are plotted except those from Sites 547, 763, and selected samples from Site 545 (see text for discussion). Disparities between nanofossil and planktic foraminiferal biostratigraphies of individual sections lead to minor inconsistencies in ages plotted against the integrated chronology (Fig. 2). Isotopic ages of mid-plate volcanic episodes are shown (dark shaded pattern refers to cluster of ages of Ontong Java and Manihiki plateaus; light shaded pattern refers to total range of ages). The key to the symbols is given in Figure 17.
Figure 22. Comparison of (A) composite data set (this study) with data sets of (B) Ingram et al. (1994). Squares are data points of samples analyzed by Ingram et al. (1994); circles are points taken from Koepnick et al. (1985) with revised biostratigraphy from Tarduno et al. (1989). Note that biostratigraphic corrections result in age errors of as much as 10 m.y.; both Ingram et al. (1994) and Koepnick et al. (1985) data sets are calibrated to Gradstein et al. (1994) time scale. (C, D) Jones et al. (1994b), (E) McArthur et al. (1994), and (F) Jenkyns et al. (1995). Solid curve in each graph (except F) shows the local-linear smooth of the data set (see Fan and Gijbels [1996] for a description of this technique); dashed curve shows the smoothed composite curve. The kernel function used for the local-linear smooth is the Gaussian function with a bandwidth of 0.75 for the data collected here, and 1.5 for the data sets of other authors. The higher bandwidth is a result of the lower sample density of previous studies. Because of the data gap in C and D, the data sets are divided in two at the gap, and two shorter curves are generated. The ages of samples in F were calculated using tie points from the composite curve (A) as follows: 97 Ma—0 m below sea floor (mbsf), 104 Ma—250 mbsf, 107 Ma—350 mbsf, 114 Ma—434.5 mbsf, 122 Ma—900 mbsf, and assuming constant sedimentation rates in between. On the basis of analyses on NBS-SRM987, we have added 0.00011 to the $^{87}\text{Sr}/^{86}\text{Sr}$ analyses of Koepnick et al. (1985) and 0.000025 to the values of Jones et al. (1994b), and subtracted 0.000025 from the values of Ingram et al. (1994). Positions of stage boundaries are shown.
123 Ma (weighted mean = 122.4 ± 0.8 Ma) (Mahoney et al., 1993; Tejada et al., 1996). Paleomagnetic data show that interbedded limestone and basalt at Ontong Java Plateau Site 807 are normally magnetized and are thought to correlate to the Cretaceous long-normal polarity interval (Tarduno et al., 1991). Combined with biostratigraphic data (Erba, 1994), this suggests an early Aptian age (between about 120.5 and 120 Ma in the Gradstein et al. [1994] time scale). The inconsistency between the biostratigraphic and isotopic ages from Ontong Java may suggest that the age of MO in the Gradstein et al. (1994) time scale is slightly too young, or that the duration of Ontong Java volcanism was longer than previous estimates (Tarduno et al., 1991). Although it is impossible to differentiate between these alternatives, we assume the full range of Ontong Java ages from biostratigraphic, paleomagnetic, and isotopic data (Fig. 21).

The Manihiki Plateau is isotopically dated in only one location (123 Ma; Mahoney et al., 1993). The large Kerguelen Plateau in the southern Indian Ocean apparently formed in a relatively short period of time, between 114 and 110 Ma (Whitechurch et al., 1992; Coffin and Eldholm, 1994), but this inference is based on isotopic ages from only four locations. Rates of ridge-crest crustal production (e.g., Kominz, 1984) are complicated by the Cretaceous long-normal polarity interval, which makes it difficult to assess relative changes in crustal production over the 38 m.y. duration of the chron, and to detect ridge jumps and plate reorganizations that, if unrecognized, artificially inflate estimates of crustal production (Heller et al., 1996). The Cretaceous long normal in the Gradstein et al. (1994) time scale begins at 120.5 Ma and ends at 83.5 Ma (Gradstein et al., 1994). This implies a shorter duration than that assumed by Larson (1991), and thus would increase somewhat the magnitude of ocean-ridge crustal production.

**Significance of Sr Isotopes to Crustal Production Estimates.** The Aptian-Albian decrease in seawater 87Sr/86Sr values is compatible with increased rates of hydrothermal activity associated with increased rates of ocean-ridge crust production (Jones et al., 1994b; Ingram et al., 1994). However, the oceanic ridges (Jones et al., 1994b; Larson, 1994) and plateaus (Ingram et al., 1994; Ingram and Richter, 1994) have both been identified as sources of increased hydrothermal Sr.

The composite seawater Sr-isotope curve offers evidence for the timing of the potentially enhanced mid-Cretaceous ocean crust production, as well as a sense of the relative magnitude of crustal production through time. The onset of accelerated crust production is well dated by the records derived from Sites 417 and 641. In both cases, the decrease in 87Sr/86Sr values starts several meters above MO (Figs. 13 and 16). If a large plume is responsible for both the long-normal polarity interval and accelerated ocean-ridge crustal production (Larson and Olson, 1991), then the 0.5 to 1 m.y. lag between the beginning of the long-normal and the onset of the decrease in seawater 87Sr/86Sr values may reflect the long (several million years) residence time of Sr as well as the time required for Sr to be leached from the new crust. If mid-plume hydrothermal activity were the sole source of volcanic Sr (Ingram et al., 1994), and most of the volcanism on the Ontong Java and Manihiki Plateaux had terminated by 120 Ma, as available data suggest (e.g., Tarduno et al., 1991; Mahoney et al., 1993), then the seawater Sr isotope decrease also lags slightly the end of the volcanic pulse (Fig. 21).

Ingram et al. (1994) interpreted their Aptian-Albian Sr-isotope data in terms of two negative excursions, each associated with the eruption of a large igneous province (Ontong Java and Kerguelen Plateaux). However, local-linear smoothing of all of the relevant data sets shows a single decrease in 87Sr/86Sr values in the Aptian-Albian stages (Fig. 22, A, C and F), not two as would be expected if the current dates for the Ontong Java, Manihiki, and Kerguelen Plateaux are representative. If volcanism ceased in most locations on Ontong Java by 121.5 Ma (Mahoney et al., 1993) and on Manihiki at 123 Ma, there is a 6 m.y. gap between volcanism on these plateaus and Kerguelen at a time when Sr-isotope ratios in the composite curve are descending rapidly (Fig. 21).

The timing of the onset of the early Aptian decrease in 87Sr/86Sr values combined with the shape of the composite Sr-curve suggest that either the current isotopic ages for the oceanic plateaus are not representative, or that leaching
of Sr from oceanic plateaus takes millions of years, or that another source, the ridge crests or a subducted plateau, supplied volcanic Sr to the oceans. Without more isotopic ages from midplate features, it is difficult to conclusively differentiate between these alternatives. However, the existing ages interpreted in light of the composite Sr-isotope curve support the notion of increased oceanic-ridge crustal production during the Aptian–early Albian stages.

Ingram et al. (1994) correlated the decrease in $^{87}$Sr/$^{86}$Sr values around the Cenomanian-Turonian boundary (93.5 Ma in Gradstein et al., 1994; Fig. 21). Low $^{87}$Sr/$^{86}$Sr values extend from the early Turonian through the Coniacian (McArthur et al., 1994), and part of this interval corresponds to the Caribbean volcanic and secondary volcanic phases on the Kerguelen and Ontong Java Plateaus between 89 and 85 Ma (Duncan et al., 1994; Bercovici and Mahoney, 1994).

The composite curve suggests that a 15 m.y. interval encompassing the middle Albian–late Cenomanian was not associated with increased volcanic Sr flux. These data conflict with Larson’s (1991) crustal production curve, which suggests that there was increased crustal production over this interval. Unfortunately, the Sr-isotope record is limited in that it cannot yet place constraints on the absolute magnitude of enhanced crustal production in the mid-Cretaceous. In addition to uncertainties in the relative contributions of midplate and ocean-ridge hydrothermal activity, the relationship between increased crustal production and increased hydrothermal Sr flux is not well established, and the response of weathering rates to any increased CO$_2$ fluxes to the atmosphere is not well understood (Kump, 1989; Berner, 1990; Caldeira and Rampino, 1991; Jones et al., 1994b).

Seawater Sr-Isotope Curves: Volcanism and Anoxic Events. Several authors have commented on possible relationships between “anoxic events,” volcanism, and the Sr-isotope record (e.g., Vogt, 1989; Jones and Jenkins, 1995). The coincidence in timing between lowered $^{87}$Sr/$^{86}$Sr values and deposition of organic-rich sediments is thought to be indirectly linked through sea level and volcanism. The close synchrony of the onset of early Aptian volcanic activity and the “Selli” anoxic event (OAE1a) (e.g., Coccioni et al., 1987) can be seen in Sr-isotope data from Site 641. Here, the onset of the decrease in $^{87}$Sr/$^{86}$Sr values leading to the Aptian trough correlates closely with an interval lacking calcareous microfossils which corresponds to the anoxic event (Bralower et al., 1994) (Fig. 16). Strontium-isotope values from Site 551 increase just below the organic-rich sediments deposited during the Cenomanian-Turonian boundary oceanic anoxic event (Bralower, 1988), but show a substantial decrease immediately above (Fig. 15). Although we have no explanation for the increase in $^{87}$Sr/$^{86}$Sr values below the black shale, we tentatively conclude that the decrease in $^{87}$Sr/$^{86}$Sr values probably resulted from a volcanic episode that led to the major early Turonian transgression (e.g., Haq et al., 1987).

SUMMARY

An investigation of mid-Cretaceous seawater Sr-isotope stratigraphy was carried out by measuring foraminifers and inorganic bivalves from 12 DSDP and ODP sections. Diagenetic alteration is identified by observing fossil preservation, measuring trace elements, and stacking $^{87}$Sr/$^{86}$Sr values of samples of similar age. The composite curve formed by samples that have undergone little or no alteration shows a trough of low $^{87}$Sr/$^{86}$Sr values in the Aptian and early Albian stages, higher but constant values for the rest of the Albian and Cenomanian stages, followed by a decrease in $^{87}$Sr/$^{86}$Sr values in the early Turonian stage. Disparities between the composite curve and published Sr-isotope stratigraphies are largely related to the different biostratigraphies and assumptions used to estimate age, rather than to diagenesis and analytical procedures. The age and shape of the trough in $^{87}$Sr/$^{86}$Sr values in the Aptian-Albian stages combined with the available isotopic age dates are consistent with some volcanic Sr being derived from increased crustal production rates at spreading centers.

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